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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 1 of 8

Complete if Known

Application Number	10/042,226
Filing Date	January 11, 2002
First Named Inventor	Bernd RIEDL et al.
Group Art Unit	1614
Examiner Name	Dwayne C. Jones
Attorney Docket Number	BAYER-0024-A

U.S. PATENT DOCUMENTS

Examiner Initials *	Cite No. ¹	U.S. Patent Document		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number	Kind Code ² (if known)			
/JDA/		4,546,191		Nishiyama et al.	10-8-1985	
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/JDA/		6,380,218		Marfat et al.	4-30-2002	
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FOREIGN PATENT DOCUMENTS

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		Office ³	Number ⁴	Kind Code ⁵ (if known)				
/JDA/		EP	0690344		Konica Corp.	1-3-1996		
/JDA/		WO	02/092576		Boehringer Ingelheim Pharm Inc.	11-21-2002		
		WO	03/062763	A2	Bayer Corp.	8-15-2002		
		WO	02/083028	A1	Boehringer Ingelheim Pharmaceuticals Inc.	10-24-2002		
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/JDA/		WO	98/20868		Tracey et al.	5-22-1998		
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		WO	02/14311		Amgen Inc.	2-21-2002		
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/JDA/		EP	0709225	B1	Nippon Paper Industries Co.	5-1-1996		

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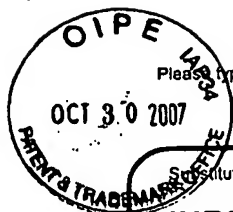
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Group Art Unit	1614
Examiner Name	Dwayne C. Jones
Attorney Docket Number	BAYER-0024-A

OTHER PRIOR ART -- NON PATENT LITERATURE DOCUMENTS

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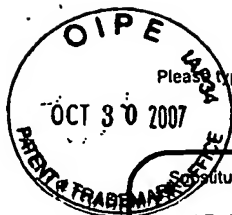
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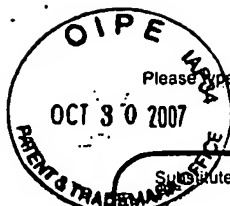
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Sheet 4 of 8

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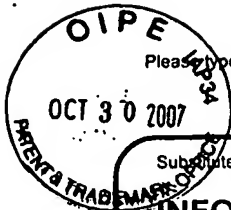
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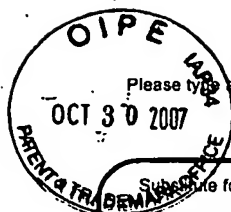
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		Application Number	10/042,226
		Filing Date	January 11, 2002
		First Named Inventor	Bernd RIEDL et al.
		Group Art Unit	1614
		Examiner Name	Dwayne C. Jones
		Attorney Docket Number	BAYER-0024-A

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Examiner Initials *	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ²
		Aucclair, et al., "BAY 43-9006 (Sorafenib) is a potent inhibitor of FLT3 tyrosine kinase signaling and proliferation in AML cells," 90 th Annual Meeting, April 16-20, 2005, Anaheim/Orange County, CA	
		Murphy et al., "BAY 43-9006 controls tumor growth through inhibition of vascular development," 96 th Annual Meeting, April 16-20, 2006, Anaheim/Orange County, CA	
		Sprengsen et al., "Novel treatment strategies in clear cell metastatic renal cell carcinoma," <i>Anti-Cancer Drugs</i> , 2005, 16:700-717	
		Thaimattam et al., "3D-QSAR-CoMFA, CoMSIA studies on substituted ureas as Raf-1 kinase inhibitors and its confirmation with structure based studies," <i>Bioorganic & Medicinal Chemistry</i> , 12(2004) 6415-6425	
		Danson et al., "Improving outcomes in advanced malignant melanoma," <i>Drugs</i> , 2005, 65(6):722-742	
		Heim et al., "Antitumor effect and potentiation or reduction in cytotoxic drug activity in human colon carcinoma cells by the Raf kinase inhibitor (RKI) BAY 43-9006," <i>International Journal of Clinical Pharmacology and Therapeutics</i> , Vol. 41, No. 12/2003 (616-617)	
/JDA/		Richly et al., "Results of a phase I trial of BAY 43-9006 in combination with doxorubicin in patients with primary hepatic cancer," <i>International Journal of Clinical Pharmacology and Therapeutics</i> , Vol. 42, No. 11/2004 (650-651)	
/JDA/		Hubbard, "Oncogenic mutations in B-Raf: some losses yield gains," Skirball Institute of Biomolecular Medicine and Department of Pharmacology, New York University School of Medicine, New York, NY; <i>Cell</i> Vol. 116, pp. 764-766, March 19, 2004	
		Thompson et al., "Recent progress in targeting the Raf/MEK/ERK pathway with inhibitors in cancer drug discovery," <i>Curr. Opin. Pharmacol.</i> , 2005 Aug., 5(4):350-6	
		Moore et al., "Phase I study to determine the safety and pharmacokinetics of the novel Raf kinase and VEGFR inhibitor BAY 43-9006, administered for 28 days on/7 days off in patients with advanced, refractory solid tumors," <i>Annals of Oncology</i> , 16:1688-1694, 2005	
		Ahmad et al., "Kinase inhibition with BAY 43-9006 in renal cell carcinoma," <i>Clinical Cancer Research</i> , Vol. 10, 6388s-6392s, 15 Sept. 2004	

Examiner Signature	/James Anderson/	Date Considered	01/17/2008
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		Clark et al., "Safety and pharmacokinetics of the dual action raf kinase and vascular endothelial growth factor receptor inhibitor, BAY 43-0006, in patients with advanced, refractory solid tumors," <i>Clin. Cancer Res.</i> , 2006;11(16), 1 August 2006, 5472-5480	
		Wilhelm et al., "BAY 43-0006: preclinical data," <i>Curr. Pharm. Des.</i> , 2002, 8(25):2255-7	
		Wright et al., "Clinical trials referral resource. Current clinical trials of BAY 43-0006, Part 1," <i>Oncology</i> , 2005 Apr, 40(4):400-502	
		Patent Abstracts of Japan, Publication No. 02-023937, published 01-26-1990	
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		Wissner et al., "Analogues of platelet activating factor. 7: Bis-aryl amide and bis-aryl urea receptor antagonists of PAF," <i>J. Med. Chem.</i> , 1992, 35, 4770-4780	
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		Escudier et al., "Randomized phase III trial of the raf kinase and VEGFR inhibitor sorafenib (BAY 43-0006) in patients with advanced renal cell carcinoma (RCC)," Meeting: 2005 ASCO Annual Meeting, Category: Genitourinary Cancer, Subcategory: Kidney Cancer, Abstract No. 4510	
		Eisen et al., "Phase I trial of BAY 43-0006 (sorafenib) combined with dacarbazine (DTIC) in metastatic melanoma patients," Meeting: 2005 ASCO Annual Meeting, Category: Melanoma, Subcategory: Melanoma, Abstract No. 7506	

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